



Office of Research Integrity-  
Human Subjects

OR I(HS)7.03



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Approved By:  
ORI Executive  
Director

\*Signature on file

Date:

Date First Effective:  
01/25/2017

Approved by:  
Biomedical Chair



UNLV Office of Research Integrity- Human Subjects and Institutional Review Board Standard Operating Procedures		
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believe that the patient may not comprehend a research protocol, the patient should not be approached for research participation. Investigators should also delay approaching patients for research participation if the patient's capacity for informed consent is impaired by medications.

Non-clinical Biomedical and Behavioral Research Involving Pregnant Women to which Subpart B may not apply

The IRB recognizes that some types of research that could be conducted with pregnant women do not fit specifically within the regulatory requirements for approval. For example, the additional safeguards require

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